

This Page Is Inserted by IFW Operations
and is not a part of the Official Record

BEST AVAILABLE IMAGES

Defective images within this document are accurate representations of the original documents submitted by the applicant.

Defects in the images may include (but are not limited to):

- BLACK BORDERS
- TEXT CUT OFF AT TOP, BOTTOM OR SIDES
- FADED TEXT
- ILLEGIBLE TEXT
- SKEWED/SLANTED IMAGES
- COLORED PHOTOS
- BLACK OR VERY BLACK AND WHITE DARK PHOTOS
- GRAY SCALE DOCUMENTS

IMAGES ARE BEST AVAILABLE COPY.

**As rescanning documents *will not* correct images,
please do not report the images to the
Image Problem Mailbox.**

12

EUROPEAN PATENT APPLICATION

21 Application number: 86102884.3

51 Int. Cl.4: **B 32 B 27/10, B 32 B 27/12,**
B 65 D 30/08

22 Date of filing: 05.03.86

30 Priority: 06.03.85 JP 44440/85

71 Applicant: **MITSUBISHI GAS CHEMICAL COMPANY, INC., 5-2, Marunouchi 2-chome Chiyoda-Ku, Tokyo (JP)**

43 Date of publication of application: 10.09.86
Bulletin 86/37

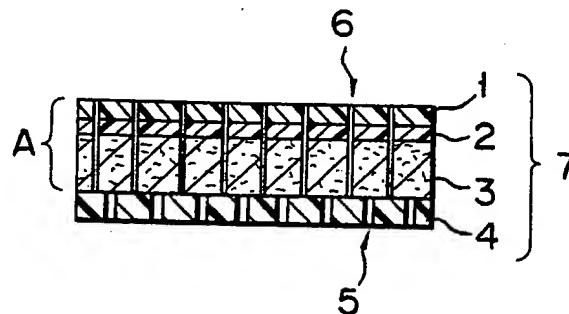
72 Inventor: **Komatsu, Toshio, 5-19-2, Kanamachi, Katsushika-ku Tokyo (JP)**
Inventor: **Takahashi, Hideyuki, 5-17-4, Nijuku, Katsushika-ku Tokyo (JP)**

84 Designated Contracting States: **DE FR GB IT NL**

74 Representative: **Sajda, Wolf Eckart, Dipl.-Phys. et al, MEISSNER, BOLTE & PARTNER Postfach 86 06 24, D-8000 München 86 (DE)**

54 **Method for manufacturing gas permeable package materials and gas permeable packaging materials obtainable thereby.**

57 A method of manufacturing a gas permeable packaging material (7), which comprises the steps of forming a laminate (A) composed of an outer plastic film (1) and a gas permeable inner sheet (3) with a first heat-sealable layer (2) interposed therebetween, perforating the laminate (A) with a number of micro-pores (6), which extend from the outer film side (1) to at least the inner sheet (3), heat-sealing a second heat-sealable film (4) with or without a number of small-pores (5) to the inner sheet (3), and, if the second heat-sealable film (4) is not porous, forming a number of small-pores in the second heat-sealable film (4), a pore ratio of the second heat-sealable film (4) being greater than that of the outer film (1).



1

5

Method for manufacturing gas permeable packaging materials
and gas permeable packaging materials obtainable thereby

10

The present invention relates to a method of manufacturing gas-permeable material for packaging a deoxidization agent, a freshness-preserving agent, and other agents and to gas permeable packaging materials.

15

Gas permeable materials have been used for packaging various agents, such as a drying agent, an insecticide, a deoxidization agent, a freshness-preserving agent, and the like. When these agents are used for preserving foods, the materials must be not only gas permeable, but also water-resistant, damage-resistant and hygienically safe.

20

The composite material consisting of a sheet of paper laminated with a perforated polyethylene film has been used for packaging a deoxidization agent. The gas permeability of the material is controlled by the ratio of the area occupied by the pores of the film to the entire surface area of the film (hereinafter called "pore ratio"). This material is, however, not resistant to water. Hence, when the bags made of the composite material and containing a deoxidization agent are used to preserve food of a high moisture content, they are damped whereby the agent oozes out through the pores. In the worst case, the bags may be broken, and the agent may come into contact with the food.

35 The composite material is not preferred from a hygienical

30

25

point of view.

The gas permeability of the polyethylene-paper material cannot always be well controlled. During the thermal bonding of the polyethylene film and the sheet of paper, the pores are deformed, whereby the material fails to have the desired gas permeability. In the bags made of such material and containing a deoxidization agent, it is difficult to control the oxygen absorption rate.

Japanese Patent Disclosures Sho 53-51096 and Sho 53-50065 disclose a packaging material consisting of two perforated plastic films laminated together and a packaging material consisting of two perforated plastic films and a sheet of paper interposed between these films. These materials are more water-resistant than a sheet of paper. Their gas permeability is, however, impaired since an adhesive such as polyethylene is used to adhere the plastic films together, or to adhere the paper sheet to the plastic films. Hence, with these materials it is practically impossible to control the gas permeability by changing the pore ratio.

Another similar composite material for packaging a deoxidization agent is known. This is a laminate made by adhering a plastic film to a layer of adhesive such as polyethylene, perforating the resultant laminate and heat-sealing laminate to a perforated heat-sealable film with a sheet of paper interposed between the laminate and heat-sealable film. This composite material has a drawback. It cannot have a desired gas permeability since the pores of the laminate and heat-sealable film are deformed during the heat sealing, inevitably changing the gas permeability, or the chips made in perforating the laminate stick in the pores, adversely reducing the gas permeability.

Japanese Patent Disclosure Sho 56-124440 discloses a packaging material made by preparing two perforated laminates each consisting of a plastic film and a sheet

1 of paper, and by adhering the paper sheets of the lami-
nates are bonded with an adhesive. It is also extremely
difficult with this packaging material to control the
gas permeability, since a layer of the adhesive such as
5 ethylene-vinyl acetate copolymer is interposed between
the paper sheets.

Japanese Patent Disclosure Sho 56-124441 also dis-
closes a packaging material comprised of two laminates.
One of the laminates consists of a plastic sheet and a
10 sheet of paper, and is perforated. The other laminate
consists of a plastic film having a gas permeability of
3000 ml/m² 24 Hr. atm. or more, and a sheet of paper.
The paper sides of these two laminates are faced and
sealed to each other with an adhesive layer interposed
15 therebetween to integrally form a packaging material.
With this packaging material it is also hard to control
the gas permeability because of the layer of an adhesive
such as ethylene-vinyl acetate copolymer.

In summary, with the conventional packaging ma-
20 terials it is extremely difficult to control the gas
permeability accurately.

It is an object of the present invention to
provide a method of manufacturing a packaging material
which is greatly water-resistant and whose gas permea-
25 bility can easily be controlled and a packaging material
having improved properties.

According to the invention, there is provided a
method of manufacturing a gas permeable packaging ma-
terial, said method comprising the steps of:

30 forming a laminate by heating an outer plastic film
and a gas permeable inner sheet, with a first heat-
sealable layer interposed between the film and sheet;

forming a large number of small-pores in the lami-
nate, extending from the outer film to at least the
35 inner sheet; and

placing, on the inner sheet of the laminate, a
second heat-sealable film having a number of small-pores
and a pore ratio greater than that of the outer plastic

film, and heating the second heat-sealable film, thereby adhering the second heat-sealable film to the inner sheet of the laminate.

According to the invention, there is provided another method of manufacturing a gas permeable packaging material, said method comprising the steps of:

forming a laminate by heating an outer plastic film and a gas permeable inner sheet, with a first heat-sealable layer interposed between the film and sheet;

forming a large number of small-pores in the laminate, extending from the outer film to at least the inner sheet;

placing, on the inner sheet of the laminate, a second heat-sealable film and heating the same, thereby adhering the second heat-sealable film to the inner sheet of the laminate; and

forming a number of small-pores in the laminate from the second heat-sealable film, extending through at least the second heat-sealable film but not through the outer film, a pore ratio of the second heat-sealable film being greater than that of the outer film.

This invention can be more fully understood from the following detailed description when taken in conjunction with the accompanying drawings, in which:

Fig. 1 is a cross-sectional view of the gas permeable packaging material manufactured by a method according to the present invention;

Fig. 2 is a cross-sectional view of part of a bag made of the packaging material shown in Fig. 1; and

Fig. 3 illustrates the relation between deoxidization time and pore ratio formed in a unit area.

A strong plastic film is preferable for the outer film of the packaging material according to this invention, since it is not broken during the manufacture of bags and during the use of the bags. The film is made of polyethylene terephthalate, polyamide, polypropylene, polycarbonate or cellophane. Of these materials,

polyethylene terephthalate or polyamide are preferable since they are strong and easy to handle in manufacturing the packaging material. It is desired that the outer film has a thickness of 50 microns or less, more preferably 10 to 20 microns. When the thickness is greater than 50 microns, the outer film will become difficult to handle.

There are no limitations to the gas permeability of the outer film, but it is generally desired that the oxygen permeability of the film be not more than 1000 ml/m² 24 Hr. atm., preferably not more than 500 ml/m² 24 Hr. atm.

When it is necessary to perform a printing on the outer film, it should be printed on the side away from the side which will contact food. This measure taken, the food will not be contaminated with the printing ink. This side of the outer film can be solid-painted after characters have been printed. In this case, the contents of the bag made of the packaging material, if oozing out of the inner film, cannot be seen from outside, whereby the outer appearance of the bag is not impaired.

The first heat-sealable film of the packaging material of the invention is used to adhere the outer film to the gas permeable inner sheet. It is a plastic film of polyethylene, ethylene-vinyl acetate copolymer, ionomer resin, polybutadiene or vinyl chloride. Alternatively, it can be made of heat sealable coating agent such as ordinary hot-melt agent, hot-melt emulsion, ionomer latex, ionomer emulsion, polyethylene emulsion or ethylene-vinyl acetate copolymer emulsion. When the first heat-sealable film is a plastic one, its thickness is preferably 10 to 70 microns, more preferably 15 to 40 microns. When the first heat-sealable film is a heat sealable coating agent, it is desired that the heat sealable coating agent is coated in an amount of 0.5 to 30 g/m², more preferably 5 to 20 g/m².

The first heat-sealable film is adhered to or

coated on the outer film which is not gas permeable. When a strong adhesion between the outer film and the gas permeable inner sheet is required, two heat-sealable films can be used.

5 Paper or nonwoven fabric is used for the gas permeable inner sheet of the packaging. The paper may be Japanese paper, rayon-mixed Japanese paper or ordinary paper. The basis weight of the paper used ranges from 15 g/m² to 100 g/m². The nonwoven fabric can be a wet-
10 processed one, a dry-processed one or a spun-bonded one. It can be made of polyamide, polyethylene terephthalate, rayon, or the like. The paper or nonwoven fabric can be coated with a water-repellant or an oil-repellant.

The second heat-sealable film, which is gas permeable, of the packaging material of the invention is
15 made of polyethylene, polyethylene-vinyl acetate copolymer, ionomer, polybutadiene or vinyl chloride.

The method according to the invention will now be described with reference to the drawings attached
20 hereto. First, as shown in Fig. 1, outer film 1, first heat-sealable film 2, and gas permeable inner sheet 3 are laid one upon another, and are then heated, thereby adhering film 1 to inner sheet 3 and providing laminate A. Laminate A can be prepared in other ways. For
25 example, film 2 is formed by applying molten material into the gap between film 1 and sheet 3, in which case the resultant laminate is passed through cold rollers after the material has solidified. Alternatively, film 2 is adhered to outer film 1, then inner sheet 3
30 is laid on film 2, and finally the three members are heated and bonded together, thus forming laminate A.

Three-layer laminate A is perforated with a needle, either hot or cold, thus forming a number of small-pores
35 6 opening in outer film 1. The diameter of pores 6 and the number of pores 6 per unit area are selected in accordance with the desired gas permeability of the packaging material, which generally ranges at most 30%

in pore ratio. To be more specific, the major axis of pores 6 may range from 0.02 to about 3 mm, in order to prevent the enclosed material from oozing out or being wetted and in view of manufacturing efficiency. These pores must be deep enough to extend from the outer side of film 1 into the surface region of inner sheet 3. The pores can be through holes, in which case it is easy to perforate laminate A.

Second heat-sealable film 4 is laid on gas permeable, inner sheet 3. The unfinished product consisting of laminate A and film 4 is heated, whereby film 4 is adhered to sheet 3. Small-pores 5 are formed in second heat-sealable film 4 and partially in inner sheet 3, either before or after the thermal adhesion. When film 4 is perforated after the thermal adhesion, pores 5 must not be so deep that they extend through inner sheet 3. It is preferred that pores 5 are made in such positions that they do not communicate with pores 6 of laminate A. If pores 5 are elliptical, their major axis may range from 0.02 mm to about 5 mm in order to prevent the enclosed material from oozing out or being wetted and in view of manufacturing efficiency. However, the size of pore 5 is not critical. They are formed in general in such numbers that film 4 have a pore ratio of 0.2 to 30%. The pore ratio of film 4 should be set such that even if the pores in film 4 are deformed by the thermal adhesion to laminate A, the pore ratio remaining in the film 4 should be larger than that of the outer film 1. For example, it is desirable that its pore ratio must be 50% or more greater than that of laminate A. Hence, the gas permeability of the packaging material according to the invention is determined by the pore ratio of film 1.

Second heat-sealable film 4 without pores is adhered to inner sheet 3, thus forming a gas permeable, packaging material 7. Film 4 can be laminated with laminate A in various methods such as thermal lamination, extrusion lamination, dry lamination and wet lamination.

Packaging material 7, thus manufactured, is folded, as shown in Fig. 2, with second heat-sealable film 4 positioned inside. The three edges of the folded are heat-sealed after deoxidization agent 8 has been placed between the halves of packaging material, whereby a bag containing deoxidization agent 8 is provided.

The bag shown in Fig. 2 can be used to contain not only a deoxidization agent, but also drying agents, insecticides, alcohol-based freshness-preserving agents, or the like. The control of the gas permeability of such a packaging material is important particularly when the material is used to manufacture bags containing deoxidization agents including sulfite, hydrogen sulfite, 2-thionite, hydroquinone, catecol, resorcinol, pyrogallol, gallic acid, ronalite, ascorbic acid, iso-ascorbic acid, and its salt, sorbose, glucose, lignin, dibutyl-hydroxytoluene, butyl-hydroxyanisole, ferrous salt, metal powder such as iron powder, a deoxidization agent producing carbon dioxide gas, and a deoxidization agent absorbing carbon dioxide gas. Among them, iron powder, ascorbic acid, iso-ascorbic acid, and its salt are most preferable. Packaging material 7 of this invention, whose gas permeability can easily be controlled, is suitable as the material for such bags.

In the method of the present invention, pores 6 formed in outer film 1, which controls the gas permeability of the packaging material, are neither deformed nor closed. Hence, the material can therefore have the desired pore ratio. Since the heat-sealable films used in this invention have pores, they do not impair the gas permeability of the packaging material. The gas permeability is determined by only the pore ratio of laminate A. It is, therefore, easy to control the gas permeability of the material. Furthermore, the material is very resistant to water.

[Examples]

The present invention will be described in more

detail with reference to examples.

Example 1

Characters were printed on one side of polyethylene terephthalate film (i.e., outer film 1) having a thickness of 12 microns. Then, this side of film 1 was coated with white ink. Film 1 and polyethylene film (i.e., first heat-sealable film 2) were subjected extrusion lamination, thereby forming a composite sheet. The composite sheet was laminated with a sheet of Japanese paper having a basis weight of 50 g/m^2 (i.e., inner sheet 3), thus providing laminate A. Pores having a diameter of 0.2 mm were formed in laminate A, at intervals of 5 mm in both the vertical direction and the horizontal direction. Sheet A thus perforated had a pore ratio of 0.12%. The perforated laminate A was adhered to a polyethylene film having a thickness of 30 microns (i.e., second heat-sealable film 4). Pores having a diameter of 0.2 mm were formed in film 4, at intervals of 2.5 mm in the vertical direction and at intervals of 1.5 mm in the horizontal direction. These pores did not penetrate inner sheet 3. Perforated film 4 had a pore ratio of 0.83%. As a result, a packaging material was obtained. The material was cut into rectangular pieces having a width of 50 mm and a length of 100 mm. These pieces were placed on a table, with second heat-sealable film 4 turned upward. An iron-based deoxidization agent is put on each piece, in an amount of 1.5 g. The piece was folded double, and the three sides of the folded piece were sealed by a three-direction bar heater, thereby providing a bag containing the iron-based deoxidization agent.

Controller 1

A controller (hereinafter called "Controller 1") was prepared. A sheet of white paper having a basic weight of 50 g/m^2 was laminated with a polyethylene film, thereby providing a packaging material. The polyethylene film had pores having a diameter of 0.2 mm and

formed at intervals of 2.5 mm in the vertical direction and at intervals of 1.5 mm in the horizontal direction. This packaging material was cut into rectangular pieces having a width of 50 mm and a length of 100 mm. These
5 pieces were placed on a table, with the polyethylene film turned upward. The same iron-based deoxidization agent is put on each piece, in an amount of 1.5 g. The piece was folded double, and the three sides of the folded piece were sealed by a three-direction bar heater, there-
10 by providing a bag containing the deoxidization agent.

The bag of Example 1 and that of Controller 1 were each put on a wet layer of cotton, the water content of which was 5 cm³/g. Each bag and the wet cotton layer were put into a KOP/PE bag and sealed therein along with
15 500 cm³ of air. The KOP/PE bags were stored at 35°C. Two days later, the oxygen concentrations in the KOP/PE bags were analyzed, and the bags were taken out of the KOP/PE bags and were examined to see the condition of the deoxidization agent. The results were as shown in the
20 following table.

Table 1

	Oxygen concentration after 2 days	Condition after 14 days	
		Exudation	Wetting
25 Example 1	Not more than 0.1%	None	None
30 Controller 1	2.1%	Rust transferred to cotton	All surface wet

Example 2

Characters were printed on one side of a poly-
amide film (i.e., outer film 1) having a thickness of
35 15 microns. Then, this side of the polyamide film was coated with white ink. The polyamide film and a polyethylene film (i.e., first heat-sealable film 2) were

laminated, thereby forming a composite sheet. The composite sheet was laminated with a sheet of white, oil-resistant paper having a basis weight of 40 g/m^2 (i.e., inner sheet 3), thus forming laminate A. Laminate A was cut
5 into smaller sheets. Pores having a diameter of 0.1 mm were formed in five of these smaller sheets, so that the five sheets had different pore ratios, i.e., 0.008%, 0.039%, 0.079%, 0.118% and 0.157%, respectively. A polyethylene film having a thickness of 30 microns (i.e.,
10 second heat-sealable film 4) was laid on the paper sheet of each perforated smaller sheet. Pores having a diameter of 0.2 mm were formed in film 4, at intervals of 2.5 mm in the vertical direction and at intervals of 1.5 mm in the horizontal direction. These pores did not
15 penetrate inner sheet 3. Perforated film 4 had a pore ratio of 0.83%. As a result, five kinds of packaging materials were obtained. These materials were cut into rectangular pieces having a width of 50 mm and a length of 100 mm. These pieces were placed on a table, with
20 second heat-sealable film 4 turned upward. An iron-based deoxidization agent was put on each piece, in an amount of 2.0 g. The piece was folded double, and the three sides of the folded piece were sealed by a three-direction bar heater, thereby providing a bag containing the iron-based deoxidization agent.
25

The bags of Example 2 were each put into a KOP/PE bag and sealed therein along with 500 cm^3 of air. The KOP/PE bags were stored at 25°C . The oxygen concentration of each KOP/PE bag was measured at 24th hour, 36th
30 hour, 48th hour, 60th hour and 72nd hour. The relation between the pore ratios of the packaging materials, on the one hand, and the oxygen concentrations measured at different times, on the other hand, was as shown in the following table:

Table 2

5	Number of micro- pores of outer material side (Number/100cm)	Pore ratio (%)	Concentration of oxygen				
			24th hr.	36th hr.	48th hr.	60th hr.	72ond hr.
	100	0.008	12.4%	9.4%	7.1%	4.9%	3.2%
	500	0.039	9.7	5.9	2.6	0.1	0.0
	1,000	0.079	5.5	0.6	0.0	-	-
10	1,500	0.118	2.0	0.0	-	-	-
	2,000	0.159	0.0	-	-	-	-

As Table 2 shows; it was confirmed that the speed
 15 of oxygen absorption of the bags was controlled by the
 pore ratio of the outer laminate.

Example 3

Characters were printed on one side of a polyethy-
 lene terephthalate film (i.e., outer film 1) having a
 20 thickness of 12 microns. Then, this side of film 1
 was coated with white ink. The polyethylene terephtha-
 late film, a polyethylene film having a thickness of
 20 microns (i.e., first heat-sealable film 2), and a
 nonwoven fabric sheet of polyethylene terephthalate
 25 having a basis weight of 35 g/m² (i.e., inner sheet 3)
 were subjected to thermal lamination, thus providing a
 laminate A. Pores having a diameter of 0.2 mm were
 formed in laminate A, at intervals of 5 mm in both the
 vertical direction and the horizontal direction. Lami-
 30 nate A thus perforated had a pore ratio of 0.12%. The
 perforated sheet A was laminated with a polyethylene
 vinyl acetate film having a thickness of 40 microns
 (i.e., second heat-sealable film 4). Pores having a
 diameter of 0.2 mm were formed in film 4, at intervals
 35 of 2.0 mm in the vertical direction and the horizontal
 direction, thus imparting to film 4 a pore ratio of
 0.78%. (These pores did not penetrate inner sheet 3.)

As a result, a packaging material was obtained. The material was cut into rectangular pieces having a width of 50 mm and a length of 100 mm. These pieces were placed on a table, with second heat-sealable film 4 turned upward. An iron-based deoxidization agent is put on each piece, in an amount of 1.5 g. The piece was folded double, and the three sides of the folded piece were sealed by a three-direction bar heater, thereby providing a bag containing the iron-based deoxidization agent. Each bag was put into a KOP/PE bag and sealed therein along with 500 g sponge cake. The KOP/PE bags were left to stand at 25°C. Two days later, the oxygen concentrations in the KOP/PE bags were found to be 0%. One month later, the sponge cake in all bags of Example 2 was well preserved, no mildewed at all. Nor did the deoxidization agent ooze out, and the outer appearance of the bags was good.

As a controller, 500 g of sponge cake was put into a KOP/PE bag, which was left to stand at 25°C. One week later, the bag was opened. The sponge cake was found to have been mildewed.

Example 4

The same experiments as in Example 1 were repeated except that the diameter of the pores 6 were varied to 0.1 mm (pore ratio, 0.21%), 0.3 mm (pore ratio, 1.88%) and 0.4 mm (pore ratio, 3.34%). As a result, oxygen concentration after two days in each cases was less than 0.1% and no exudation nor wetting was seen after 14 days of storing at 35°C.

Claims:

1. A method of manufacturing a gas permeable packaging material, which is characterized by comprising the steps of:

forming a laminate (A) by heating an outer plastic
5 film (1) and a gas permeable inner sheet (3), with a first heat-sealable layer (2) interposed between the film (1) and sheet (3);

forming a large number of small-pores (6) in the laminate (A), extending from the outer film (1) to at
10 least the inner sheet (3); and

placing, on the inner sheet (3) of the laminate, a second heat-sealable film (4) having a number of small-pores (5) and a pore ratio greater than that of the outer plastic (1) film, and heating the second heat-sealable film (4), thereby adhering the second heat-sealable film (4) to the inner sheet of the laminate (A).
15

2. A method of manufacturing a gas permeable packaging material, which is characterized by comprising the steps of:

forming a laminate (A) by heating an outer plastic
20 film (1) and a gas permeable inner sheet (3), with a first heat-sealable layer (2) interposed between the film (1) and sheet (3);

forming a large number of small-pores (6) in the laminate (A), extending from the outer film (1) to at
25 least the inner sheet (3);

placing, on the inner sheet (3) of the laminate (A), a second heat-sealable film (4) and heating the same, thereby adhering the second heat-sealable film (4) to
30 the inner sheet of the laminate (A); and

forming a number of small-pores (5) in the laminate (A) from the second heat-sealable film (4), extending through at least the second heat-sealable film (4) but not through the outer film (1), a pore ratio of the
35 second heat-sealable film (4) being greater than that

1 of the outer film (1).

3. The method according to claim 1 or 2, characterized in that the thickness of the outer film (1) is not more than 50 microns.

5 4. The method according to any of claims 1 to 3, characterized in that said heat-sealable films (2, 4) are made of a thermoplastic material.

5. The method according to any of claims 1 to 4, characterized in that said heat-sealable films (2, 4)
10 are formed by coating a heat sealable coating agent on the gas permeable inner sheet (3) in an amount of 0.5 to 30 g/m².

6. The method according to any of claims 1 to 5, characterized in that said gas permeable inner sheet (3) is
15 made of paper or nonwoven fabric.

7. The method according to claim 6, characterized in that said paper or nonwoven fabric has a basis weight of 15 to 100 g/m².

8. The method according to any of claims 1 to 7,
20 characterized in that said laminate (A) is formed by adhering three layers by heat-pressing.

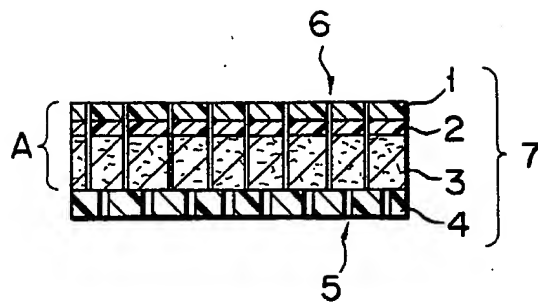
9. The method according to any of claims 1 to 7, characterized in that said laminate (A) is formed by pouring a molten heat sealable coating agent into the gap
25 between said outer film (1) and said gas permeable inner sheet (3) and then cooling the heat sealable coating agent.

10. The method according to any of claims 2 to 9, characterized in that said second heat-sealable film (4) is laminated with said laminate (A) by heat-pressing.

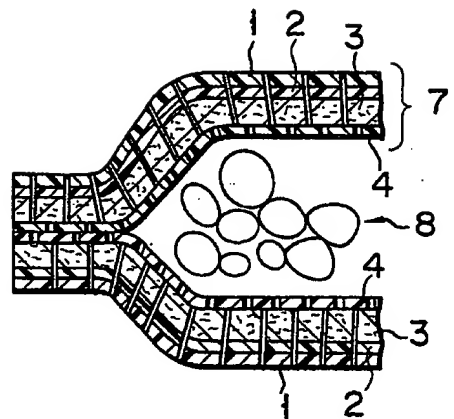
30 11. The method according to any of claims 1 to 10, characterized in that said second heat-sealable film (4) has a pore ratio of 0.2 to 30 %, the pore ratio of said second heat-sealable film (4) being greater than that of said outer film (1).

35 12. A gas permeable packaging material obtainable by a method according to any of claims 1 to 11.

F I G. 1

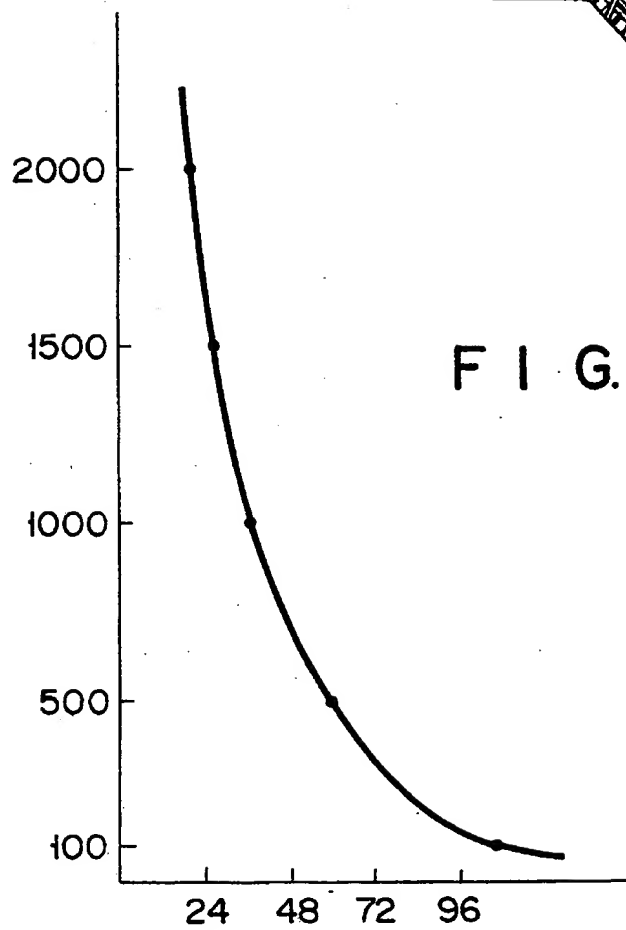


F I G. 2



PORE RATIO

F I G. 3



DEOXIDATION TIME (Hr.)